

PCT

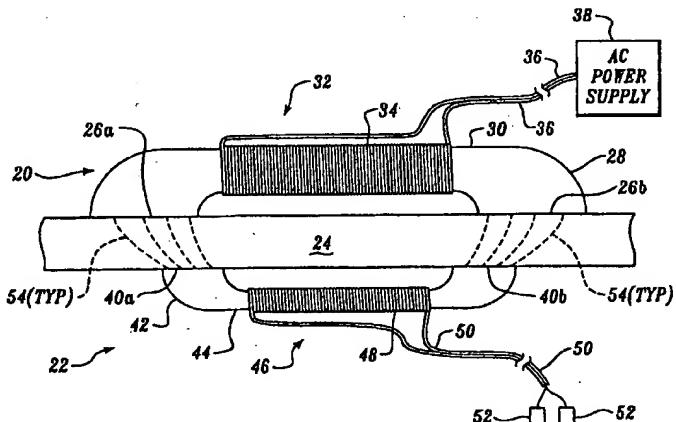
WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : A61N 2/00, 2/02	A1	(11) International Publication Number: WO 98/24510 (43) International Publication Date: 11 June 1998 (11.06.98)
(21) International Application Number: PCT/US97/17391		(81) Designated States: AU, CA, JP, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).
(22) International Filing Date: 26 September 1997 (26.09.97)		
(30) Priority Data: 08/756,945 2 December 1996 (02.12.96) US		Published With international search report. With amended claims.
(71) Applicant: LIGHT SCIENCES LIMITED PARTNERSHIP [US/US]; 1065 - 12th Avenue N.W. #E5, Issaquah, WA 98027 (US).		
(72) Inventors: CHEN, James, C.; 2011 - 87th Place N.E., Bellevue, WA 98004 (US). WILKERSON, Brian, D.; 4850 - 156th Avenue N.E. #87, Redmond, WA 98052 (US). HUSTON, Darrin; 37330 - 246th Avenue S.E., Enumclaw, WA 98022 (US). BROWN, David, J.; 3240 Porter Street, Enumclaw, WA 98022 (US).		
(74) Agent: ANDERSON, Ronald, M.; Law Offices of Ronald M. Anderson, Suite 1710, 500 - 108th Avenue N.E., Bellevue, WA 98004 (US).		

(54) Title: ELECTROMAGNETIC COIL CONFIGURATIONS FOR POWER TRANSMISSION THROUGH TISSUE



(57) Abstract

A system for improved transcutaneous power transfer efficiency comprises a transmitter coil (20) applied to the outer surface of a cutaneous layer (24) on a patient's body by adhesive tape or other appropriate supporting material, and a receiver coil (22) implanted inside the patient's body. The transmitter coil and receiver coil include cores (28, 42) having pole faces (26a, 26b, 40a, 40b) with a substantially larger area than the cross section of the core at other locations. In addition, the core of the receiver coil (42) is substantially shorter than that of the transmitter coil (28) so that the lines of flux (54) produced by the transmitter coil tend to pass through the pole faces (40a, 40b) of the receiver coil in greater density than they would if the pole faces of the transmitter and receiver cores were spaced identically. The transmitter core (28) forms an angled portion adjacent each of the pole faces so that the main portion of the transmitter core (28) is disposed generally parallel, and closely spaced to the surface of the cutaneous layer (24) against which the transmitter core (28) is placed.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

ELECTROMAGNETIC COIL CONFIGURATIONS FOR POWER TRANSMISSION THROUGH TISSUE

Field of the Invention

The present invention is generally directed to devices for 5 electromagnetically conveying power across an interface, and more specifically, to transmitter and receiver coils that are used to convey power transcutaneously to a medical device implanted within a body of a patient.

Background of the Invention

The use of implanted medical devices such as pacemakers, artificial hearts, 10 and nerve stimulators has become increasingly more prevalent with the continuing development of medical technology. These types of devices are usually energized with an electrical current supplied from a rechargeable implanted battery or from an external source. While it is possible to directly connect an external power supply to an implanted medical device via leads that extend transcutaneously from 15 the patient's body, the risk of infection and other complications makes it preferable to couple power to the implanted devices without passing leads through the skin. One approach that is used provides for supplying power to the implanted device through a radio frequency (RF) energy transfer from an external transmitter. The RF signal produced by the transmitter is transmitted through the 20 skin to an RF receiver that is connected to a power supply on the implanted device. Typically, the received RF signal must be converted to a direct current (DC) signal that is employed to power the implanted device. However, the energy transfer efficiency of such systems is relatively poor across larger gaps.

An alternative approach for transcutaneous power transmission employs an 25 electromagnetic transmitter coil that is energized with a relatively low frequency (e.g., 60 Hz) alternating current (AC) signal, producing a magnetic field that induces a corresponding current in an implanted electromagnetic receiver coil. For example, "C-shaped" half toroidal core transmitter and receiver coils usable to

transmit power transcutaneously are disclosed in commonly assigned U.S. Patent application, Serial No. 08/451,831 (allowed) and set to issue as U.S. Patent No. 5,571,152 and U.S. Patent application, Serial No. 08/705,334, filed August 29, 1996. While the configuration of a C-shaped core having a plurality of 5 turns of conductor wound about its central portion that is shown in these applications provides for relatively good power transfer, it would clearly be preferable to develop transmitter and receiver coils that are more efficient and comparatively more compact than those disclosed in these earlier references.

The C-shaped core used for the coils in these prior application has two 10 problems. First, the transmitter core tends to extend outwardly too far from the skin's surface, making it more difficult to attach and mount the transmitter coil in a proper disposition, for example, at a site on the chest of a patient. The mass and profile of the C-shaped core cause it to extend outwardly from the surface of the skin. It is contemplated that the core might be secured in place with adhesive tape 15 of other suitable binding material during the transfer of power to the internal receiving coil. However, the C-shaped coil extends outwardly sufficiently far that it would likely be difficult to maintain the pole faces of the core against the skin surface as required for efficient transcutaneous power transfer. In addition, the receiving coil should also be relatively compact, and should have a low profile to 20 minimize the internal volume required to position it adjacent the dermal layer in the patient's body. Secondly, it is important that the transmitter and receiver coils tolerate at least some misalignment between their pole faces, since it may not be possible to determine the exact position of an implanted receiving coil when 25 mounting the transmitter coil to the skin of the patient. In addition, it is desirable that the configuration and size of the core of the transmitting coil and of the receiving coil be optimized for the transcutaneous power transfer to occur over the required distances. Most conventional transformers are designed to couple over relatively short gaps between their respective pole faces, but the gap between the pole faces of the transmitter coil core and the receiver coil core used for 30 transcutaneous power transmission may be several centimeters.

Summary of the Invention

In accord with the present invention, an electromagnetic coil configuration for conveying power transcutaneously is defined that includes a core of 35 magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section. The intermediate section has a smaller cross-sectional size than that of the first and second pole faces to provide an increased

5 pole face area for conveying a transcutaneous magnetic flux. A plurality of turns of an electrical conductor are wound around the intermediate section. This electrical conductor carries an induced current if the coil is used as a receiver that is electromagnetically excited by an external source of the transcutaneous magnetic flux. Alternatively, if the coil is used as a transmitter of the transcutaneous magnetic flux, the turns of the electrical conductor carry a varying electrical current supplied by a power supply to which the electrical conductor is adapted to be connected.

10 In one embodiment, the intermediate section comprises a first section and a second section that are respectively coupled to the first pole face and the second pole face. The first section has a first winding comprising a portion of the plurality of the turns of the electrical conductor, and the second section has a second winding comprising another portion of the plurality of the turns of the electrical conductor. In one configuration, the first section is directly coupled to 15 the second section, and the core is generally V-shaped. In another configuration, the intermediate section further comprises a third section that extends between the first section and the second section, so that the core is generally U-shaped. Preferably, the first and second sections include an angled portion disposed adjacent to the first and second pole faces, enabling the first and second sections 20 to extend generally to one side of the first and second pole faces, so that the core has a low profile relative to the first and second pole faces.

25 In another embodiment, the intermediate section includes a mid-portion that is elongate and flattened, and which extends in a direction generally parallel to the first and second pole faces. The plurality of turns of the electrical conductor are wound around the mid-portion.

It is also preferable that the core comprise a plurality of layers of the magnetically permeable material, adjacent layers of the magnetically permeable material being separated by a layer of a dielectric material, to minimize losses in the core due to eddy currents. The same result is achieved if the core comprises a 30 plurality of discrete elements such as particles or rods that are formed of the magnetically permeable material, which are bonded together with a dielectric material.

35 Another aspect of the present invention is directed to a system that includes a transmitter coil and a receiver coil for transcutaneously transferring power. A cross-sectional size of the first and second pole faces of the transmitter substantially differs from the cross-sectional size of the first and second pole faces

of the receiver. Preferably, the transmitter core is longer than the receiver core and an area of the first and second pole faces of the transmitter core is substantially greater than that of the first and second pole faces of the receiver core.

5

Brief Description of the Drawing Figures

The foregoing aspects and many of the attendant advantages of this invention will become more readily appreciated as the same becomes better understood by reference to the following detailed description, when taken in conjunction with the accompanying drawings, wherein:

10

FIGURE 1 is a side view of first embodiments of a transmitter coil and a receiver coil for transcutaneously coupling power in accord with the present invention;

FIGURE 2 is a plan view of the first embodiment of the transmitter coil, which is substantially similar in appearance to the receiver coil;

15

FIGURE 3 is a side elevational view of a patient in which a medical device has been implanted, showing the first embodiment of the transmitter coil positioned on the patient's torso to couple power transcutaneously to a receiver coil (not shown) within the patient's body;

20

FIGURE 4 is a side view of a second embodiment of a transmitter coil and the receiver coil of FIGURE 1, disposed on opposite sides of a tissue layer;

FIGURE 5 is an isometric view of a third embodiment of a transmitter or receiver coil;

FIGURE 6 is an isometric view of a fourth embodiment of a transmitter or receiver coil;

25

FIGURE 7 is an isometric view of a fifth embodiment of a transmitter or receiver coil;

FIGURE 8 is a cut-away view of a cross section through a core of a transmitter or receiver coil that includes a plurality of longitudinally extending stacked layers of a magnetically permeable material alternating with a dielectric material;

FIGURE 9 is a cut-away view of a cross section through a core of a transmitter or receiver coil that includes a plurality of concentric layers of magnetically permeable material alternating with a dielectric material;

35

FIGURE 10 is a cut-away view of a cross section through a core of a transmitter or receiver coil that includes a plurality of rods of magnetically permeable material bound in a matrix of a dielectric material;

FIGURE 11 is a cut-away view of a cross section through a core of a transmitter or receiver coil that includes a plurality of particles of magnetically permeable material bound in a matrix of a dielectric material; and

5 FIGURE 12 is a cut-away view of a cross section through a core of a transmitter or receiver coil comprising helically rolled sheets of a magnetically permeable material and a dielectric material.

Description of the Preferred Embodiments

Transmission and receiver coils previously developed for transcutaneous power transmission to energize a probe to administer photodynamic therapy (PDT) have included a "C-shaped" core around which was wound a helical coil of an electrical conductor. The transmitter coil in this system is adapted to be coupled to an external power supply that provides a 60 Hz AC. When the electrical current flows through the windings of the transmitter coil, power is transmitted over a distance of several centimeters to a similarly shaped receiver 10 coil that is coupled to a plurality of light emitting diodes (LEDs) in an implantable PDT probe. Ongoing efforts to develop more efficient designs for the transmitter 15 and receiver coils have resulted in the present invention.

FIGURE 1 shows a first embodiment of the present invention in which a transmitter coil 20 is electromagnetically coupled to a receiver coil 22 to convey 20 power transcutaneously through an intervening cutaneous layer 24. In this embodiment of the invention, core faces 26a and 26b of a core 28 have a surface area that is substantially larger than a cross-sectional area of the remaining portion of the core. The larger surface area results from flaring the ends of core 28 outwardly proximate core faces 26a and 26b. FIGURE 2 illustrates the relative 25 size of core faces 26a and 26b compared to the remainder of core 28.

Core 28 includes an intermediate section 30 that connects the two portions of the core on which core faces 26a and 26b are disposed. A transmitter winding 32 comprises a plurality of turns of an electrical conductor 34, preferably in two or more layers of helical windings, since the flux density produced by 30 transmitter coil 20 is directly proportional to the number of turns of electrical conductor 34 that are wrapped around intermediate section 30. A lead 36 extends from transmitter winding 32 to an AC power supply 38, that would likely be connected to an AC line. It is also contemplated that AC power supply 38 can include an external battery pack and an inverter (neither shown). Also, the current 35 supplied through lead 36 can have a sinusoidal, pulse, or other time varying

waveform. Furthermore, in the preferred embodiment, the frequency of the electrical current supplied to transmitter coil 20 is preferably less than 500 Hz, since the electromagnetic coupling between transmitter coil 20 and receiver coil 22 has been found to be more efficient at the typical spacing between the 5 transmitter and receiver coil at such frequencies. Another advantage of using a relatively low frequency electrical current is that radio frequency shielding around the transmitter coil is not required.

Receiver coil 22 is generally similar to transmitter coil 20, except that it is relatively more compact and shorter in length. Pole faces 40a and 40b on the ends 10 of a core 42 of receiver coil 22 are also relatively larger in area than other portions of the core and are formed by flaring out the ends of core 42 proximate to the pole faces. An intermediate section 44 couples the portions of core 42 on which pole faces 40a and 40b are disposed. Wrapped around intermediate section 44 is a receiver winding 46 comprising a plurality of turns of an electrical conductor 48; 15 preferably in two or more layers. Electrical conductor 48 is connected through a lead 50 to terminals 52 that are adapted for coupling the receiver coil to a medical device (not shown), which is implanted within the body of the patient, so that electrical current induced to flow within receiver winding 46 can be used to energize the medical device.

20 Further details of cores 28 and 42 are discussed below, in connection with various configurations used to avoid losses due to eddy currents circulating therein. Preferably, cores 28 and 42 are fabricated using a material that has a relatively high magnetic permeability, such as μ metal or other alloys typically used for electromagnetic cores. It has been found that enhanced performance is 25 obtained for coupling power transcutaneously between transmitter coil 20 and receiver coil 22 by making the length of core 42 sufficiently shorter than that of core 28 so that for a given thickness of cutaneous layer 24, lines of electromagnetic flux 54 tend to flow between pole faces 26a and 26b in a direction passing through corresponding pole faces 40a and 40b of the receiver coil. To 30 optimize power transfer between the coils, the spacing between core faces 26a and 26b should be substantially greater than the thickness of cutaneous layer 24. Previously it was thought that the length of core 28 should equal that of core 42, making the distance between the center of the pole faces of the transmitter coil equal to the distance between the center of the pole faces of the transmitter coil. 35 However, it has been determined that by making the distance between the center of the pole faces of the receiver coil shorter than that between the centers of the

pole faces of the transmitter coil, the flux density passing through the pole faces of the receiver coil is increased, thereby enhancing the efficiency of the transcutaneous power transfer process.

Referring now to FIGURE 3, a patient 60 is schematically illustrated to show how transmitter coil 20 is applied to a torso 62 of the patient for transcutaneous power transfer to a receiver (not shown), which is implanted immediately opposite transmitter coil 20, inside the patient's body. If transmitter coil 20 is intended to be used while patient 60 is mobile, the external power source (not shown in this Figure) could be attached to a belt worn by the patient and would include a storage battery. Furthermore, transmitter coil 20 can be adhesively secured to torso 62 of the patient to facilitate the transcutaneous power transfer required to energize an internal implanted medical device that is coupled to the receiver coil. Because of the relatively low profile of transmitter coil 20, it can be covered by loose garments so that its use by patient 60 is not evident to others. The patient can then enjoy the freedom of pursuing normal activities while the implanted medical device provides its intended therapeutic benefit.

FIGURE 4 illustrates a second embodiment of a transmitter coil 70. Transmitter coil 70 includes a core 74 having pole faces 72a and 72b disposed at each end. Core 74 extends upright above the respective pole faces and includes an intermediate section 76 that is generally transverse to the upright portions of the core connecting the two sections on which are disposed transmitter windings 78a and 78b. These transmitter windings each comprise a plurality of turns of an electrical conductor 80, preferably in multiple layers. Interconnecting electrical conductors 80' convey electrical current between transmitter windings 78a and 78b, and a lead 82 is provided for connecting the transmitter coil to a power supply (like power supply 38, shown in FIGURE 1). Core 74 can be formed as a contiguous assembly with intermediate section 76, or alternatively, the two upright portions and the intermediate section can be fabricated separately and connected together to provide a continuous flux path coupling transmitter windings 78a and 78b so that the magnetic flux produced by the flow of an electrical current through electrical conductor 80 flows through intermediate section 76 of core 74. The transmitter windings 78a and 78b around the two upright portions of core 74 include a plurality of turns of electrical conductor 80 wrapped in a plurality of layers. The two transmitter windings are coupled together by leads 80'. It should be clear that the direction of the transmitter

winding turns about each portion of core 74 is such that the magnetic flux they produce reinforces rather than cancels in core 74.

As indicated in FIGURE 4, the overall distance between the center of pole faces 72a and 72b is substantially greater than that between the centers of pole faces 40a and 40b of receiver coil 22. This relationship between the relative lengths of the transmitter coil and receiver coil cores, as described above, provides improved efficiency for the transcutaneous power transfer between the two devices. Also, the distance between pole faces 72a and 72b should again be substantially greater than the thickness of cutaneous layer 24.

One of the problems with the design for transmitter coil 70 as shown in FIGURE 4 is its relative height Y' above the surface of cutaneous layer 24. As noted above, it is contemplated that the transmitter coil will be attached to the outer surface of cutaneous layer 24 using adhesive tape or other suitable binding material (not shown). Further, it was noted above that it would be desirable to easily hide a transmitter coil under loose clothing. However, due to the relative height of transmitter coil 70 compared to that of the much more compact transmitter coil 20, it will be apparent that securing transmitter coil 70 to patient 60 so that the transmitter coil remains upright with pole faces 72a and 72b in continuous contact with the surface of cutaneous layer 24 would be more difficult, since the transmitter coil would be more likely to shift and lean away from its desired upright position relative to the outer surface of the cutaneous layer against which it is affixed. Moreover, transmitter coil 70, which may have a height (dimension "Y") that is more than two inches, is far too evident and difficult to hide, even under loose clothing, because of the distance that it extends above the cutaneous layer.

A solution to this problem is shown in FIGURE 5, which illustrates a transmitter coil 90. In this embodiment, pole faces 92a and 92b are disposed at opposite ends of a core 94. However, unlike core 74 of transmitter coil 70, core 94 includes two angled portions 96 adjacent each end; at angled portions 96, the core turns through an angle of approximately 90° so that a plane through the upper portion of the core is generally parallel to pole faces 92a and 92b. Core 94 also includes an intermediate section 98 joining the two portions of the core about which transmitter windings 100 are formed from turns of an electrical conductor 102. As in the previous embodiments, the transmitter windings around each of these portions of core 94 include a plurality of turns of electrical conductor 102 in a plurality of layers. The two transmitter windings are coupled

together by leads 102'. Again, the direction of the transmitter winding turns about each portion of core 94 is such that the magnetic flux they produce reinforces rather than cancels in core 94. Since most of core 94 is disposed relatively close to the plane of core faces 92a and 92b, transmitter coil 90 can more readily be
5 affixed to the surface of the cutaneous layer through which electromagnetic power is transferred to a corresponding receiving coil, such as receiving coil 22. In addition, transmitter coil 90 will be less evident under the clothing of a patient, since it does not extend very high above the surface of the cutaneous layer.

10 Alternative configurations of the embodiment just discussed above are shown in FIGURES 6 and 7. In FIGURE 6, a transmitter coil 90' includes an intermediate section 98', which is relatively shorter than intermediate section 98 of transmitter coil 90. Carrying this modification to its next logical step, a transmitter coil 90" in FIGURE 7 is generally V-shaped, so that intermediate section 98" simply joins the two portions of a core 94" on which transmitter
15 windings 100 are disposed.

20 Turning now to FIGURES 8-12, a plurality of configurations for the core of the transmission coil and/or receiver coil in each of the above-described embodiments are illustrated. In FIGURE 8, a core configuration 110 comprises alternating layers 112 of a μ metal or other alloy of the type conventionally used in electromagnetic coils, and layers 114 of a dielectric material. The dielectric material in layers 114 separates the material of layers 112, which has a relatively high magnetic permeability, minimizing eddy currents that cause substantial loss in solid cores of a material that has a high magnetic permeability.

25 In FIGURE 9, a core configuration 118 includes alternating concentric layers 120 and 122. Layers 120 comprise the material having a relatively high magnetic permeability, such as μ metal or other alloy conventionally used in fabricating electromagnetic cores, and layers 122 comprise a dielectric material, such as various types of polymers characterized by having a relatively high dielectric constant.

30 FIGURE 10 shows a core configuration 130 in which a plurality of generally parallel, elongate rods 132 extend through the core in spaced-apart array, bonded in that matrix configuration by a dielectric material 134. As noted above, dielectric material 134 preferably comprises a polymer or other suitable dielectric material. Rods 132 may be formed of μ metal or other alloy
35 conventionally used to fabricate electromagnetic cores.

In FIGURE 11, a core configuration 140 comprises a plurality of spaced-apart particles 142 of a material having a relatively high magnetic permeability, such as μ metal, formed as a matrix bound by a material 144 that has a relatively high dielectric property. Once again, a polymer is preferably used for the dielectric material in this configuration.

Finally in FIGURE 12, a further alternative core configuration 150 is shown. Core configuration 150 comprises a flexible sheet 152 of a material having a relatively high magnetic permeability, such as μ metal, that is helically rolled with a flexible sheet 154 of a material having a relatively high dielectric property. An appropriate flexible polymer is preferably used for the dielectric material in this embodiment.

Although the present invention has been described in connection with several preferred forms of practicing it, those of ordinary skill in the art will understand that many other modifications can be made thereto within the scope of the claims that follow. Accordingly, it is not intended that the scope of the invention in any way be limited by the above description, but instead be determined entirely by reference to the claims that follow.

The invention in which an exclusive right is claimed is defined by the following:

1. An electromagnetic coil configuration for conveying power transcutaneously, comprising:

(a) a core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, a cross-sectional area of said intermediate section being substantially less than that of said first and second pole faces to provide an increased pole face area for conveying a transcutaneous magnetic flux; and

(b) a plurality of turns of an electrical conductor wound around said intermediate section, said electrical conductor carrying an induced current if the coil is used as a receiver that is electromagnetically excited by an external source of the transcutaneous magnetic flux, and carrying a varying electrical current supplied by a power supply to which the electrical conductor is adapted to be connected if the coil is used as a transmitter of the transcutaneous magnetic flux.

2. The electromagnetic coil configuration of Claim 1, wherein the intermediate section comprises a first section and a second section that are respectively coupled to the first pole face and the second pole face, said first section having a first winding comprising a portion of the plurality of the turns of the electrical conductor, and said second section having a second winding comprising another portion of the plurality of the turns of the electrical conductor.

3. The electromagnetic coil configuration of Claim 2, wherein the first section is connected to the second section, and the core is generally V-shaped.

4. The electromagnetic coil configuration of Claim 2, wherein the intermediate section further comprises a third section that extends between the first section and the second section, so that the core is generally U-shaped.

5. The electromagnetic coil configuration of Claim 2, wherein the first and second sections include an angled portion disposed adjacent to the first and second pole faces, said first and second sections extending generally to one side of the first and second pole faces, so that said core has a low profile relative to the first and second pole faces.

6. The electromagnetic coil configuration of Claim 1, wherein the intermediate section includes a mid-portion that extends in a direction generally parallel to the first and second pole faces, said plurality of turns of the electrical conductor being wound around said mid-portion.

7. The electromagnetic coil configuration of Claim 1, wherein the core comprises a plurality of layers of the magnetically permeable material, adjacent layers of the magnetically permeable material being separated by a layer of a dielectric material, to minimize losses in the core due to eddy currents.

8. The electromagnetic coil configuration of Claim 1, wherein the core comprises a plurality of discrete elements formed of the magnetically permeable material that are bonded with a dielectric material, to minimize losses in the core due to eddy currents.

9. The electromagnetic coil configuration of Claim 8, wherein the discrete elements comprise elongate rods that extend through the core.

10. A system for transferring power transcutaneously to energize an implanted medical device within a patient's body, comprising:

(a) a transmitter coil adapted to be disposed external to the patient's body, said transmitter coil including:

(i) a transmitter core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section; and
(ii) a plurality of turns of an electrical conductor wound around said intermediate section;

(b) a receiver coil adapted to be disposed subdermally, inside the patient's body, said receiver coil including:

(i) a receiver core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said first and second pole faces having a substantially different area than the first and second pole faces of the transmitter coil; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section and having ends adapted to couple to the medical device; and

(c) a power supply adapted to couple to the turns of the electrical conductor wound around the intermediate section of the transmitter core, said power

supply supplying a time varying electrical current to energize the transmitter coil, producing an electromagnetic field that couples transcutaneously with the receiver coil, causing a corresponding electrical current to flow in the receiver coil for energizing the medical device implanted inside the patient's body.

11. The system of Claim 10, wherein the first and second pole faces of the receiver coil have a substantially smaller area than that of the first and second pole faces of the transmitter coil.

12. The system of Claim 10, wherein the intermediate section of the transmitter coil has a cross-sectional area that is substantially smaller than the area of either of the first and second pole faces of the transmitter coil.

13. The system of Claim 10, wherein the intermediate section of the receiver coil has a cross-sectional area that is substantially smaller than the area of either of the first and second pole faces of the receiver coil.

14. The system of Claim 10, wherein the receiver coil is substantially shorter than the transmitter coil.

15. The system of Claim 10, wherein a space separating the first and second pole faces of the transmitter coil is substantially greater than a thickness of a cutaneous layer of the patient's body that is disposed between the transmitter coil and the receiver coil.

16. The system of Claim 10, wherein the intermediate section of the transmitter core comprises a first section and a second section that respectively terminate in the first pole face and the second pole face of the transmitter core, said first section having a first winding comprising a portion of the plurality of the turns of the electrical conductor, and said second section having a second winding comprising another portion of the plurality of the turns of the electrical conductor that is wound around the intermediate section of the transmitter core.

17. The system of Claim 16, wherein the first section is directly coupled to the second section, and the transmitter core is generally V-shaped.

18. The system of Claim 10, wherein the intermediate sections of at least one of the transmitter coil and the receiver coil includes a mid-portion that is

elongate and flattened, and which extends in a direction that is generally parallel to the first and second pole faces of said at least one of the transmitter coil and the receiver coil, said plurality of turns of the electrical conductor for said at least one of the transmitter coil and the receiver coil being wound around said mid-portion thereof.

19. The system of Claim 10, wherein at least one of the transmitter core and the receiver core comprises a plurality of discrete elements formed of the magnetically permeable material that are separated by a dielectric material, to minimize losses in said at least one of the transmitter core and the receiver core, due to eddy currents.

20. A system for transferring power transcutaneously to energize an implanted medical device within a patient's body, comprising:

(a) a transmitter coil adapted to be disposed external to the patient's body, said transmitter coil including:

(i) a transmitter core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said transmitter core increasing in cross-sectional size so that said first and second pole faces have an increased pole face area for conveying a transcutaneous magnetic flux; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section;

(b) a receiver coil adapted to be disposed subdermally, inside the patient's body, said receiver coil including:

(i) a receiver core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said receiver core being substantially shorter than the transmitter core and increasing in size so that said first and second pole faces of the receiver core have an increased pole face area for receiving a transcutaneous magnetic flux generated by the transmitter coil; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section and having ends adapted to couple to the medical device; and

(c) a power supply adapted to couple to the turns of the electrical conductor wound around the intermediate section of the transmitter

core, said power supply supplying a time varying electrical current to energize the transmitter coil, producing an electromagnetic field that couples transcutaneously with the receiver coil, causing a corresponding electrical current to flow in the receiver coil for energizing the medical device implanted inside the patient's body.

21. The system of Claim 20, wherein the intermediate section of the transmitter core comprises a first section and a second section that respectively terminate in the first pole face and the second pole face of the transmitter core, said first section having a first winding comprising a portion of the plurality of the turns of the electrical conductor, and said second section having a second winding comprising another portion of the plurality of the turns of the electrical conductor that is wound around the intermediate section of the transmitter core.

22. The system of Claim 21, wherein the first section is directly coupled to the second section, and the transmitter core is generally V-shaped.

23. The system of Claim 21, wherein the intermediate section of the transmitter core further comprises a third section that extends between the first section and the second section, so that the transmitter core is generally U-shaped.

24. The system of Claim 21, wherein the first and second sections of the transmitter core include angled portions disposed adjacent to the first and second pole faces of the transmitter core that turn the transmitter core to one side of the first and second pole faces, said first and second sections thereby extending generally to said one side of the first and second pole faces, so that said transmitter core has a low profile relative to the first and second pole faces of the transmitter core.

25. The system of Claim 20, wherein the intermediate sections of at least one of the transmitter coil and the receiver coil includes a mid-portion that is elongate and flattened, and which extends in a direction that is generally parallel to the first and second pole faces of said at least one of the transmitter coil and the receiver coil, said plurality of turns of the electrical conductor for said at least one of the transmitter coil and the receiver coil being wound around said mid-portion thereof.

26. The system of Claim 20, wherein at least one of the transmitter core and receiver core comprises a plurality of layers of the magnetically permeable material, adjacent layers of the magnetically permeable material being separated by a layer of a dielectric material, to minimize losses in said at least one of the transmitter core and receiver core due to eddy currents.

27. The system of Claim 20, wherein at least one of the transmitter core and the receiver core comprises a plurality of discrete elements formed of the magnetically permeable material that are bonded together with a dielectric material, to minimize losses in said at least one of the transmitter core and the receiver core, due to eddy currents.

28. The system of Claim 27, wherein the discrete elements comprise elongate rods that extend through said at least one of the transmitter core and the receiver core.

29. The system of Claim 20, wherein the power supply provides a periodically varying electrical current to energize the transmitter coil.

30. The system of Claim 29, wherein the electrical current periodically varies at a frequency less than 500 Hz.

AMENDED CLAIMS

[received by the International Bureau on 09 January 1998 (09.01.98);
original claims 1,3,10-14,17,20,22 and 30 amended; new claims 31-34 added;
remaining claims unchanged (6 pages)]

The invention in which an exclusive right is claimed is defined by the following:

1. An electromagnetic coil configuration for conveying power transcutaneously, comprising:
 - (a) a core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, a transverse cross-sectional area of said intermediate section being substantially less than a transverse cross-sectional area of said first and second pole faces to provide an increased pole face area for conveying a transcutaneous magnetic flux; and
 - (b) a plurality of turns of an electrical conductor wound around said intermediate section, said electrical conductor carrying an induced current if the coil is used as a receiver that is electromagnetically excited by an external source of the transcutaneous magnetic flux, and carrying a varying electrical current supplied by a power supply to which the electrical conductor is adapted to be connected if the coil is used as a transmitter of the transcutaneous magnetic flux.
2. The electromagnetic coil configuration of Claim 1, wherein the intermediate section comprises a first section and a second section that are respectively coupled to the first pole face and the second pole face, said first section having a first winding comprising a portion of the plurality of the turns of the electrical conductor, and said second section having a second winding comprising another portion of the plurality of the turns of the electrical conductor.
3. The electromagnetic coil configuration of Claim 2, wherein the first section is connected to the second section, and the core has a general V-shape.
4. The electromagnetic coil configuration of Claim 2, wherein the intermediate section further comprises a third section that extends between the first section and the second section, so that the core is generally U-shaped.
5. The electromagnetic coil configuration of Claim 2, wherein the first and second sections include an angled portion disposed adjacent to the first and second pole faces, said first and second sections extending generally to one side of the first and second pole faces, so that said core has a low profile relative to the first and second pole faces.
6. The electromagnetic coil configuration of Claim 1, wherein the intermediate section includes a mid-portion that extends in a direction generally parallel to the first and second pole faces, said plurality of turns of the electrical conductor being wound around said mid-portion.
7. The electromagnetic coil configuration of Claim 1, wherein the core comprises a plurality of layers of the magnetically permeable material, adjacent layers of the magnetically permeable material being separated by a layer of a dielectric material, to minimize losses in the core due to eddy currents.

-18-

8. The electromagnetic coil configuration of Claim 1, wherein the core comprises a plurality of discrete elements formed of the magnetically permeable material that are bonded with a dielectric material, to minimize losses in the core due to eddy currents.

9. The electromagnetic coil configuration of Claim 8, wherein the discrete elements comprise elongate rods that extend through the core.

10. A system for transferring power transcutaneously to energize an implanted medical device within a patient's body, comprising:

(a) a transmitter coil adapted to be disposed external to the patient's body, said transmitter coil including:

(i) a transmitter core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section;

(b) a receiver coil adapted to be disposed subdermally, inside the patient's body, said receiver coil including:

(i) a receiver core of a substantially solid and magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said first and second pole faces having a substantially different transverse cross-sectional area than the first and second pole faces of the transmitter coil; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section and having ends adapted to couple to the medical device; and

(c) a power supply adapted to couple to the turns of the electrical conductor wound around the intermediate section of the transmitter core, said power supply supplying a time varying electrical current to energize the transmitter coil, producing an electromagnetic field that couples transcutaneously with the receiver coil, causing a corresponding electrical current to flow in the receiver coil for energizing the medical device implanted inside the patient's body.

11. The system of Claim 10, wherein the first and second pole faces of the receiver coil have a substantially smaller transverse cross-sectional area than that of the first and second pole faces of the transmitter coil.

12. The system of Claim 10, wherein the intermediate section of the transmitter coil has a transverse cross-sectional area that is substantially smaller than the transverse cross-sectional area of either of the first and second pole faces of the transmitter coil.

13. The system of Claim 10, wherein the intermediate section of the receiver coil has a transverse cross-sectional area that is substantially smaller than the transverse cross-sectional area of either of the first and second pole faces of the receiver coil.

14. The system of Claim 10, wherein the receiver coil has a length substantially shorter than that of the transmitter coil, said shorter length causing a distance between

centers of the first and second pole faces of said receiver coil to be less than a distance between centers of the first and second pole faces of said transmitter coil.

15. The system of Claim 10, wherein a space separating the first and second pole faces of the transmitter coil is substantially greater than a thickness of a cutaneous layer of the patient's body that is disposed between the transmitter coil and the receiver coil.

16. The system of Claim 10, wherein the intermediate section of the transmitter core comprises a first section and a second section that respectively terminate in the first pole face and the second pole face of the transmitter core, said first section having a first winding comprising a portion of the plurality of the turns of the electrical conductor, and said second section having a second winding comprising another portion of the plurality of the turns of the electrical conductor that is wound around the intermediate section of the transmitter core.

17. The system of Claim 16, wherein the first section is directly coupled to the second section, and the transmitter core has a general V-shape.

18. The system of Claim 10, wherein the intermediate sections of at least one of the transmitter coil and the receiver coil includes a mid-portion that is elongate and flattened, and which extends in a direction that is generally parallel to the first and second pole faces of said at least one of the transmitter coil and the receiver coil, said plurality of turns of the electrical conductor for said at least one of the transmitter coil and the receiver coil being wound around said mid-portion thereof.

19. The system of Claim 10, wherein at least one of the transmitter core and the receiver core comprises a plurality of discrete elements formed of the magnetically permeable material that are separated by a dielectric material, to minimize losses in said at least one of the transmitter core and the receiver core, due to eddy currents.

20. A system for transferring power transcutaneously to energize an implanted medical device within a patient's body, comprising:

(a) a transmitter coil adapted to be disposed external to the patient's body, said transmitter coil including:

(i) a transmitter core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said transmitter core increasing in transverse cross-sectional size so that said first and second pole faces have an increased pole face area for conveying a transcutaneous magnetic flux; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section;

(b) a receiver coil adapted to be disposed subdermally, inside the patient's body, said receiver coil including:

(i) a receiver core of a substantially solid and magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said receiver core increasing in transverse cross-sectional size so that said first and second pole

faces of the receiver core have an increased pole face area for receiving a transcutaneous magnetic flux generated by the transmitter coil; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section and having ends adapted to couple to the medical device; and

(c) a power supply adapted to couple to the turns of the electrical conductor wound around the intermediate section of the transmitter core, said power supply supplying a time varying electrical current to energize the transmitter coil, producing an electromagnetic field that couples transcutaneously with the receiver coil, causing a corresponding electrical current to flow in the receiver coil for energizing the medical device implanted inside the patient's body.

21. The system of Claim 20, wherein the intermediate section of the transmitter core comprises a first section and a second section that respectively terminate in the first pole face and the second pole face of the transmitter core, said first section having a first winding comprising a portion of the plurality of the turns of the electrical conductor, and said second section having a second winding comprising another portion of the plurality of the turns of the electrical conductor that is wound around the intermediate section of the transmitter core.

22. The system of Claim 21, wherein the first section is directly coupled to the second section, and the transmitter core has a general V-shape.

23. The system of Claim 21, wherein the intermediate section of the transmitter core further comprises a third section that extends between the first section and the second section, so that the transmitter core is generally U-shaped.

24. The system of Claim 21, wherein the first and second sections of the transmitter core include angled portions disposed adjacent to the first and second pole faces of the transmitter core that turn the transmitter core to one side of the first and second pole faces, said first and second sections thereby extending generally to said one side of the first and second pole faces, so that said transmitter core has a low profile relative to the first and second pole faces of the transmitter core.

25. The system of Claim 20, wherein the intermediate sections of at least one of the transmitter coil and the receiver coil includes a mid-portion that is elongate and flattened, and which extends in a direction that is generally parallel to the first and second pole faces of said at least one of the transmitter coil and the receiver coil, said plurality of turns of the electrical conductor for said at least one of the transmitter coil and the receiver coil being wound around said mid-portion thereof.

26. The system of Claim 20, wherein at least one of the transmitter core and receiver core comprises a plurality of layers of the magnetically permeable material, adjacent layers of the magnetically permeable material being separated by a layer of a dielectric material, to minimize losses in said at least one of the transmitter core and receiver core due to eddy currents.

27. The system of Claim 20, wherein at least one of the transmitter core and the receiver core comprises a plurality of discrete elements formed of the magnetically permeable material that are bonded together with a dielectric material, to minimize losses in said at least one of the transmitter core and the receiver core, due to eddy currents.

28. The system of Claim 27, wherein the discrete elements comprise elongate rods that extend through said at least one of the transmitter core and the receiver core.

29. The system of Claim 20, wherein the power supply provides a periodically varying electrical current to energize the transmitter coil.

30. The system of Claim 29, wherein the periodically varying electrical current has a frequency less than 500 Hz.

31. The system of Claim 20, wherein the receiver coil has a length that is substantially shorter than another length of the transmitter coil, the shorter length of said receiver coil causing a distance between centers of the first and second pole faces of said receiver coil to be less than a distance between centers of the first and second pole faces of said transmitter coil.

32. An electromagnetic coil configuration for conveying power transcutaneously, comprising:

(a) a core of a magnetically permeable material, said core being V-shaped and having a first pole face coupled to a second pole face by an intermediate section, a transverse cross-sectional area of said intermediate section being substantially less than a transverse cross-sectional area of said first and second pole faces to provide an increased pole face area for conveying a transcutaneous magnetic flux; and

(b) a plurality of turns of an electrical conductor wound around said intermediate section, said electrical conductor carrying an induced current if the coil is used as a receiver that is electromagnetically excited by an external source of the transcutaneous magnetic flux, and carrying a varying electrical current supplied by a power supply to which the electrical conductor is adapted to be connected if the coil is used as a transmitter of the transcutaneous magnetic flux.

33. A system for transferring power transcutaneously to energize an implanted medical device within a patient's body, comprising:

(a) a transmitter coil adapted to be disposed external to the patient's body, said transmitter coil including:

(i) a transmitter core of a magnetically permeable material, said transmitter core being V-shaped and having a first pole face coupled to a second pole face by an intermediate section; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section;

(b) a receiver coil adapted to be disposed subdermally, inside the patient's body, said receiver coil including:

(i) a receiver core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said first and second pole faces having a substantially different transverse cross-sectional area than the first and second pole faces of the transmitter coil; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section and having ends adapted to couple to the medical device; and

(c) a power supply adapted to couple to the turns of the electrical conductor wound around the intermediate section of the transmitter core, said power supply supplying a time varying electrical current to energize the transmitter coil, producing an electromagnetic field that couples transcutaneously with the receiver coil, causing a corresponding electrical current to flow in the receiver coil for energizing the medical device implanted inside the patient's body.

34. A system for transferring power transcutaneously to energize an implanted medical device within a patient's body, comprising:

(a) a transmitter coil adapted to be disposed external to the patient's body, said transmitter coil including:

(i) a transmitter core of a magnetically permeable material, said transmitter core being V-shaped and having a first pole face coupled to a second pole face by an intermediate section, said transmitter core increasing in transverse cross-sectional size so that said first and second pole faces have an increased pole face area for conveying a transcutaneous magnetic flux; and

(ii) a plurality of turns of an electrical conductor wound around said intermediate section;

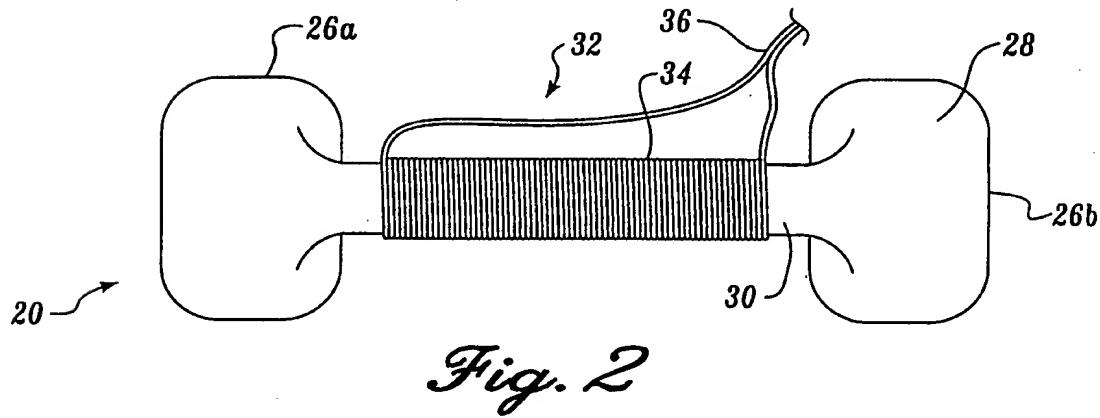
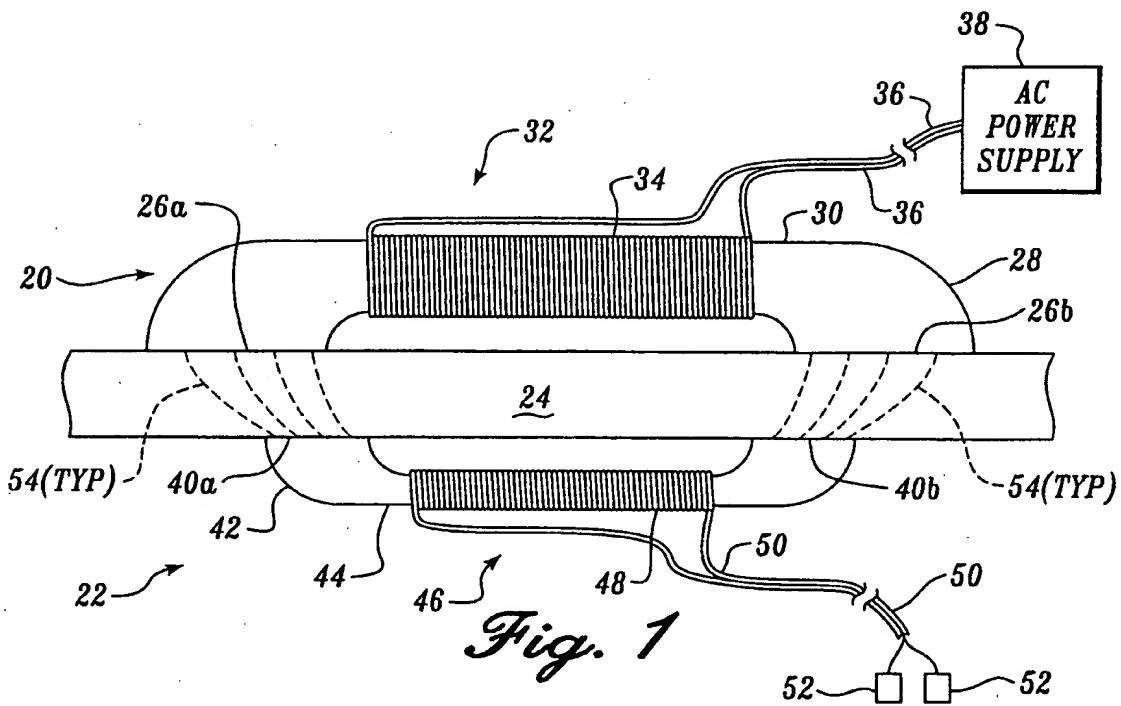
(b) a receiver coil and adapted to be disposed subdermally, inside the patient's body, said receiver coil including:

(i) a receiver core of a magnetically permeable material having a first pole face coupled to a second pole face by an intermediate section, said receiver core increasing in transverse cross-sectional size so that said first and second pole faces of the receiver core have an increased pole face area for receiving a transcutaneous magnetic flux generated by the transmitter coil; and

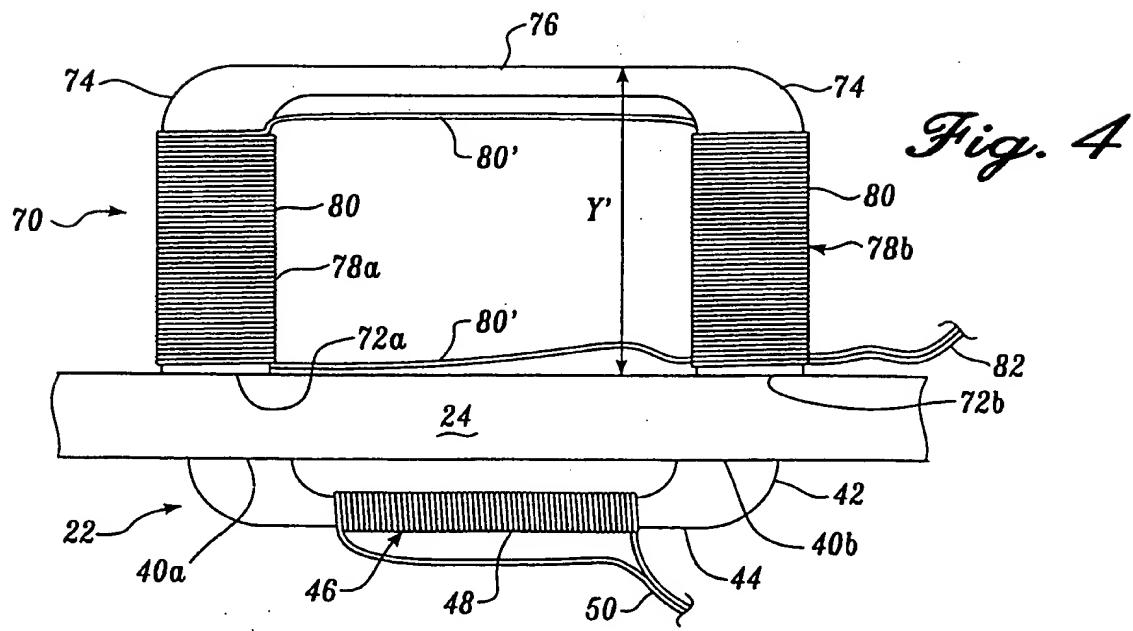
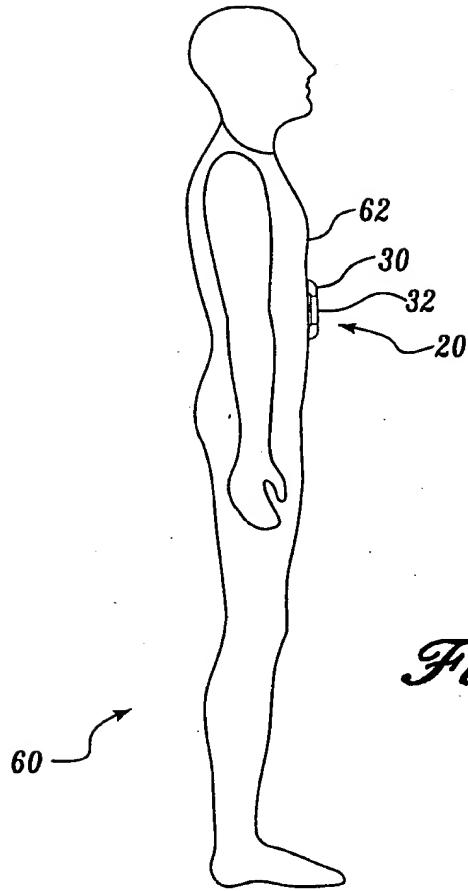
(ii) a plurality of turns of an electrical conductor wound around said intermediate section and having ends adapted to couple to the medical device; and

(c) a power supply adapted to couple to the turns of the electrical conductor wound around the intermediate section of the transmitter core, said power supply supplying a time varying electrical current to energize the transmitter coil, producing an electromagnetic field that couples transcutaneously with the receiver coil, causing a corresponding electrical current to flow in the receiver coil for energizing the medical device implanted inside the patient's body.

1/4



2/4



3/4

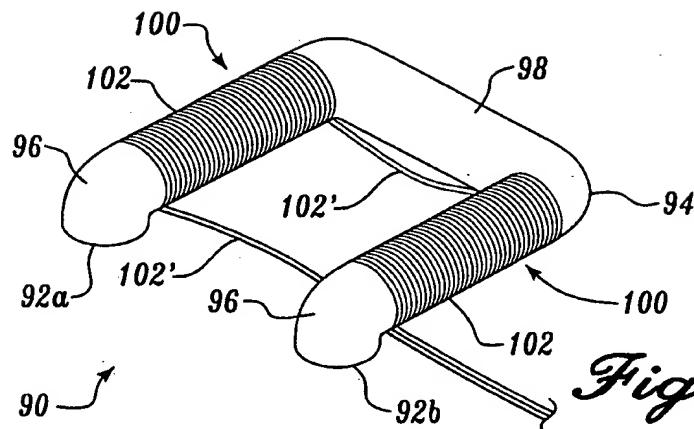


Fig. 5

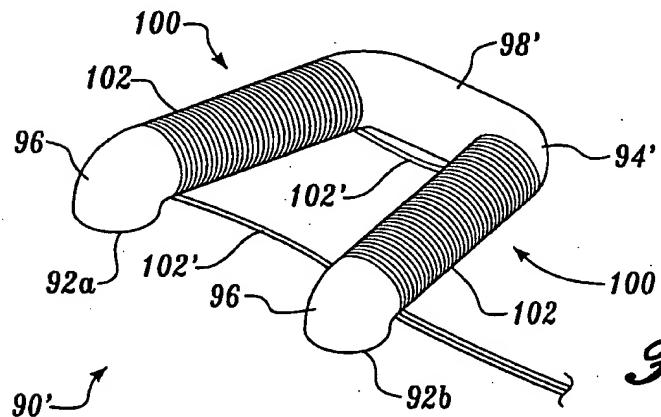


Fig. 6

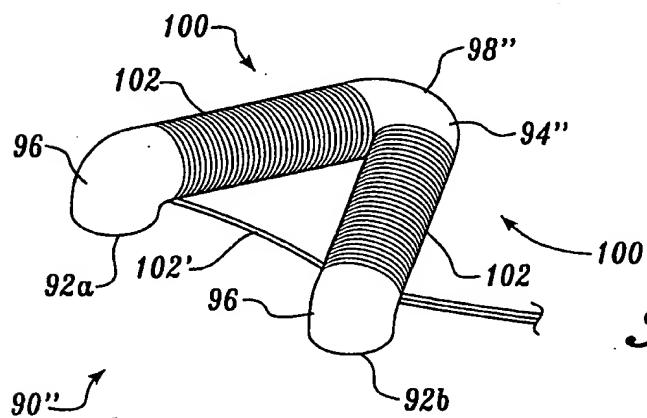


Fig. 7

4/4

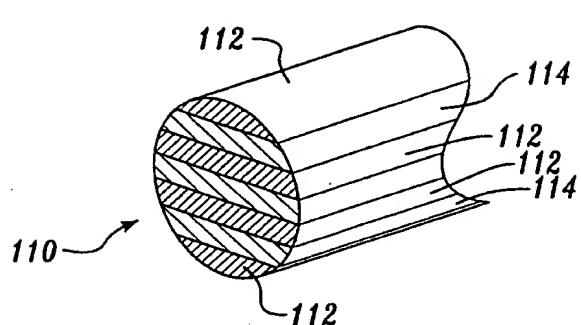


Fig. 8

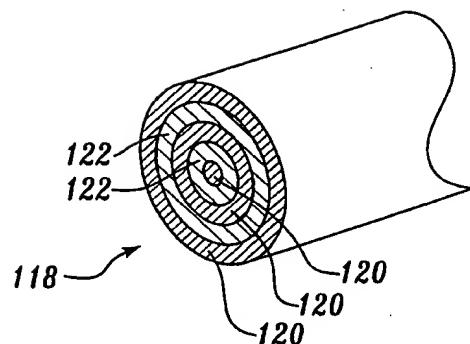


Fig. 9

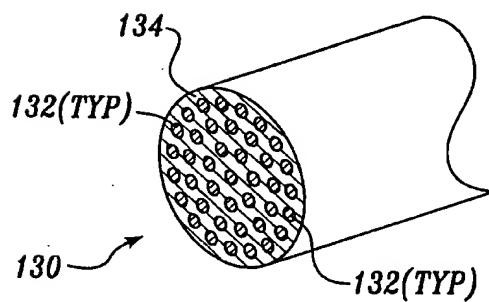


Fig. 10

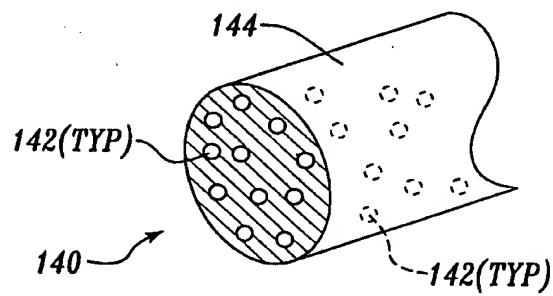


Fig. 11

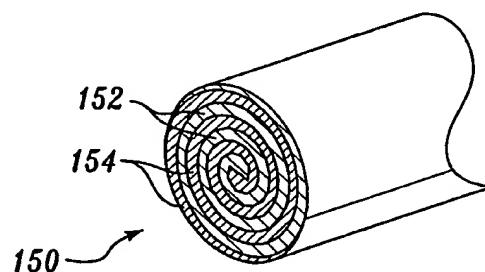


Fig. 12

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/17391

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :A61N 2/00, 02
US CL :607/2, 33, 61, 65

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 607/2, 33, 61, 65

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,000,178 A (GRIFFITH) 19 March 1991, Figs.14 and 15.	1, 2, 5
Y	US 5,314,453 A (JEUTTER) 24 May 1994, col.3, lines 38-43.	10, 11, 14, 20, 29
Y	SU 602195 A (BOREIKO V F) April 1978, translated page 1, Abstract, and figure.	10, 11, 14, 20, 29
Y	MILLER et al., Development of an Autotuned Transcutaneous Energy Transfer System, ASAIO Journal 1993, pages M706-M710.	10, 11, 14, 20, 29

 Further documents are listed in the continuation of Box C. See patent family annex.

•	Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E"	earlier document published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O"	document referring to an oral disclosure, use, exhibition or other means		
"P"	document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

05 NOVEMBER 1997

Date of mailing of the international search report

12 DEC 1997

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized Officer

CARL H. LAYNO

Telephone No. (703) 308-3694